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**Carbon fibril mfr. - obt'd. by decomposition of carbon source in contact
with metal catalyst on inorganic substrate**

Patent Assignee: HYPERION CATALYSIS INT (HYPE-N); HYPERION CATALYSIS INT
INC (HYPE-N)

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Number of Countries: 032 Number of Patents: 020

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WO 9007023	A	19900628				199029	B
AU 9049473	A	19900710				199039	
ZA 8909615	A	19900926	ZA 899615	A	19891215	199044	
EP 451208	A	19911016	EP 90901950	A	19891218	199142	
BR 8907845	A	19911008				199145	
NO 9102313	A	19910805				199145	
CA 2005642	A	19900616				199213	
JP 4504445	W	19920806	WO 89US5666	A	19891218	199238	
			JP 90502421	A	19891218		
AU 642401	B	19931021	AU 9049473	A	19891218	199349	
IL 92717	A	19940227	IL 92717	A	19891215	199419	
EP 451208	A4	19920422	EP 90901950	A		199521	
US 5500200	A	19960319	US 84678701	A	19841206	199617	
			US 86871675	A	19860606		
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			US 88149573	A	19880128		
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			US 94284755	A	19940802		
			US 95479864	A	19950607		
JP 2899407	B2	19990602	WO 89US5666	A	19891218	199927	
			JP 90502421	A	19891218		
JP 11256430	A	19990921	JP 90502421	A	19891218	199950	
			JP 9910094	A	19891218		
EP 969127	A2	20000105	EP 90901950	A	19891218	200006	
			EP 99114233	A	19891218		
KR 137224	B1	19980428	WO 89US5666	A	19891218	200013	
			KR 90701780	A	19900816		
EP 451208	B1	20000322	WO 89US5666	A	19891218	200019	
			EP 90901950	A	19891218		
			EP 99114233	A	19891218		
DE 68929243	E	20000921	DE 629243	A	19891218	200055	
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			EP 90901950	A	19891218		
CA 2005642	C	20001024	CA 2005642	A	19891215	200059	
US 6375917	B1	20020423	US 84678701	A	19841206	200232	
			US 86871675	A	19860606		
			US 86871676	A	19860606		
			US 86872215	A	19860606		
			US 88149573	A	19880128		
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			US 91746065	A	19910812		

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JP 11256430	A		7	D01F-009/127	Div ex application JP 90502421
EP 969127	A2 E			D01F-009/127	Div ex application EP 90901950 Div ex patent EP 451208
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Abstract (Basic): WO 9007023 A

Carbon source (I) is contacted at 500-1500 deg.C with an inorganic substrate (II) of size upto 400 microns supporting a multivalent transition metal present on (II) as a plurality of discontinuous catalytic sites of 35-700 Angstrom size and recovering the graphitic carbon fibril prod. (I) is e.g. methane, benzene, MeOH, CO etc. (I) can

be Al₂O₃, SiO₂, spinels, MgO etc. Atmos. used is pref. non oxidising. Metal catalyst is e.g. Fe, Mo, Co, Pt etc. Process may be carried out in an ebulliating bed. Fibril prods. may have hollow cores.

USE/ADVANTAGE - Process could also be used to mfr. silicon nitride or carbide fibrils. Fibrils have high surface area, tensile strength and elasticity modulus. Prod. is obtd. at high yield and in a thin form. Post prodn. graphitisation is not needed. Typical fibrils have 35-70 narcometer dia. and length to dia. ratios upto 100:1. Yields obtd. are above 30 times wt. of transition metal in the catalyst against a factor of 2 in the prior art. (24pp Dwg.No.0/0)

Abstract (Equivalent): US 5500200 A

The prodn. is of carbon fibrils by decomposing a source of carbon at elevated temps. in contact with a multivalent transition metal and recovering the fibrils formed.

The prodn. comprises injecting catalyst particles having a size of up to 400 mum and comprising at least one multivalent transition metal on a particulate substrate into a reactor heated to a temp. below the thermal decomposition temp. of the source of carbon, and continuously recovering the fibrils formed thereby,

said metal being present on the substrate as a multiplicity of discontinuous catalytic sites which, at least during fibril formation, have a size of about 35 - 700 A which size is measured by measuring the size of transition metal particles recovered along with produced fibrils and the fibrils comprising an inner core region of less ordered carbon atoms surrounded by an outer region of ordered carbon atoms.

Dwg.0/0

Title Terms: CARBON; FIBRIL; MANUFACTURE; OBTAIN; DECOMPOSE; CARBON; SOURCE ; CONTACT; METAL; CATALYST; INORGANIC; SUBSTRATE

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03 B114 B720 B730 B732 B833 C107 C800 C802 C803 C804 C806 C807 M411 M720 M903 M904 N441 N480 N514 N515 Q453 R042 R03124-P

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(54) **A CATALYTIC VAPOR GROWTH METHOD FOR PRODUCING CARBON FIBRILS**

EIN KATALYTISCHES DAMPFZUCHTVERFAHREN ZUR HERSTELLUNG VON
KOHLENSTOFFFIBRILLEN

PROCEDE DE CRISTALLISATION EN PHASE VAPEUR PAR VOIE CATALYTIQUE POUR LA
PRODUCTION DE FIBRILLES DE CARBONE

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(56) References cited:
EP-A- 0 198 558 WO-A-87/07559
JP-A- 57 117 622 US-A- 4 014 980
US-A- 4 435 367 US-A- 4 518 575
US-A- 4 650 657 US-A- 4 663 230
US-A- 4 767 737 US-A- 4 816 289

• E.B.Naumann: "Chemical Reactor Design"; John
Wiley & Sons, New York(US);p.xxi-xxx, 355-362
(1987)

Remarks:

The file contains technical information submitted
after the application was filed and not included in
this specification

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Description

[0001] This application refers to fibrils. It more particularly refers to carbon/graphite fibrils and to an improved process for producing such. Carbon fibrils as used herein means graphitic fibrils having high surface area, high young's modulus of elasticity and high tensile strength which are grown catalytically from available sources of carbon.

Background of the Invention

[0002] It has been known for some time that one could make fibrils by decomposing various carbon contributing molecules, such as light hydrocarbons, in contact with a suitable metal catalyst, such as for example iron alone or in combination with other metals. PCT Patent Publication WO 87/07599 to Tennent for example, relates to the production of graphitic carbon fibrils.

[0003] More particularly the invention refers to an improved process for producing such carbon/graphite fibrils. Fibrils are made according to this invention in a high temperature, catalytic process. The fibril can be made of a variety of materials, e.g. carbon, silicon nitride, silicon carbide, etc. Such fibrils have the atoms in their composition relatively ordered at their outer surfaces as they are made by this process. Thus, it can be said that this process preferably directly produces a product having a relatively crystalline outer region for substantial portions of its length and may have inner regions where its atoms are less ordered. It may, and often does, even have a hollow region axially positioned along substantial portions of its length.

[0004] Fibrils prepared by a process according to this invention are characterized by small diameters, e.g. 3.5 to 70 nanometers and high L/D up to about 100 and even more. Where the preferred structure described above is produced, it is suitably produced directly in the fibril forming process without further processing being required.

[0005] According to the present invention, Carbon fibrils can be produced in quite high yields. In this embodiment, a suitable source of carbon may be a hydrocarbonaceous material illustrated by: methane, ethane, propane, butane, benzene, cyclohexane, butene, isobutene, ethylene, propylene, acetylene, toluene, xylene, cumene, ethyl benzene, naphthalene, phenanthrene, anthracene, formaldehyde, acetaldehyde, acetone, methanol, ethanol, carbon monoxide, (a non-hydrocarbonaceous material) other similar materials, and mixtures of two (2) or more thereof. Such feed is contacted with a suitable, catalyst at elevated, fibril forming temperatures for a time sufficient to cause graphitic carbon fibrils to grow.

[0006] It is within the scope of this invention to provide a non-hydrocarbonaceous gas along with the carbon contributing reactant. Such gas might for example be hydrogen or carbon monoxide. Inert diluents are also

suitable.

[0007] The temperature of the process of this invention can vary widely depending upon the nature of the carbon source being used, however, the temperature is kept below the thermal decomposition temperature thereof. In the case of using a mixture of such carbon sources, the operating temperature should be maintained below the thermal decomposition temperature of the most temperature-sensitive carbon source in the system. Temperatures in the range of 500 to 1500°C may be found to be generally usable, depending on the carbon source used, preferably between about 600 and 900°C.

[0008] Subatmospheric, atmospheric and/or super atmospheric pressures may be used as dictated by other processing considerations. It has been found that it is desirable to provide the carbon source in the vapor state, and thus, the pressure should not be so high as to cause the carbon source to be in the liquid state under fibril forming temperature conditions. Further, it is desirable although not essential to provide a suitable gaseous diluent, such as hydrogen or inert gases, for example, nitrogen.

[0009] It is preferred that the system as a whole be non-oxidizing wherefor preferably avoiding the presence of oxygen if practical. Small amounts of these materials can be tolerated. It should be understood that the existence of oxidizing conditions, at the elevated temperatures operative for this process, will cause oxidation of the carbon source and therefor reduce the amount of carbon from such source which is available for conversion into fibrils as desired.

[0010] It may be desirable to provide suitable heat to this reaction system where and when needed. Temperature of different parts of the reactor zone may be suitably controlled to different temperatures and this is easily accomplished by using electrical resistance heating. However in larger scale industrial practice, electric resistance heating may sometimes be economically replaced by direct heating, such as for example by burning some of the carbon contributing feed to raise the temperature of the remainder of the feed, or by feeding the catalyst or the carbon contributing feed, or the diluent into the system at a sufficiently elevated temperature such that direct heat exchange of the component with each other will cause the fibril forming reaction to proceed as desired.

[0011] The nature of the catalyst seems to have a significant effect upon the yield of fibrils produced according to this invention. It is known to use iron group metals such as iron, cobalt or nickel to catalyze the conversion of carbon contributing compounds to fibrils, and such metals are within the scope of this invention. In addition, many other multivalent transition metals, including lanthanides, appear to be operative. Particularly useful catalytic metals include inter alia: iron, molybdenum, cobalt, nickel, platinum, palladium, vanadium, and chromium. The present process uses catalyst particles com-

prising as the multivalent metal, iron mixed with at least one other transition metal.

[0012] Particularly useful combinations include iron and molybdenum, iron and chromium, copper and nickel, iron and platinum, iron and tin, iron and nickel, iron and manganese, and iron and cerium.

[0013] The yield of fibrils produced according to the practice of this invention appears to be related to the physical state of the catalyst used to produce such. According to the invention, the multivalent transition metal fibril forming catalyst is present on a suitable substrate as relatively discrete catalytic sites, each 3.5 to 70 nm (35 to 700 Å), preferably 6 to 30 nm (60 to 300 Å) in size during fibril formation. These relatively discrete catalytic sites are produced by suitably applying the transition metal (in an appropriate state) to a substrate, suitably an inorganic substrate material which can include carbon/graphite.

[0014] The size of the substrate particle is a matter of some importance dependent upon the engineering or the process itself. For example, if the fibril formation is to take place in a fluid bed type of reaction zone, the substrate particle size will suitably be less than about 400 µm. If the fluid bed is an ebullient bed of catalyst particles, particle sizes of about 50 to 300 µm have been found to be preferable. If the fluid bed is an ebullient bed of fibrils containing small amounts of catalyst particles, i.e. up to about ten percent, these should preferably have a size of about 1 to 100 µm. If the fluid bed is a transport bed, either up flow or down flow, the catalyst carrying particles will suitably be less than about 10 µm preferably less than about one µm.

[0015] It has been found that depositing transition metals according to present claim 1 on small particle substrates produces a catalyst well suited to use in this invention. The substrate is a material which can conveniently withstand the rigors of fibril formation conditions, e.g. temperatures of 500 to 1500°C. Suitable substrates include carbon, graphite, inorganic oxides, etc. The particular substrate will be matched to the particular transition metal(s) catalyst such that the metal is bound strongly enough to retard migration and agglomeration but not so strongly as to prevent or retard the transition metal from catalyzing fibril formation. Illustrative, inorganic oxides include alumina, silica, magnesia, silicates, aluminates, spinels etc. Mixtures can be used.

[0016] Thus, very small particle iron such as might be produced by decomposition of iron compounds, can be deposited on very small particle alumina, e.g. fumed alumina having particle sizes of no larger than about 149 µm (100 mesh). These alumina particles may be made up of individual crystallites which are on the order of about 5 to 20 nm (50 to 200 Å), which agglomerate to form particles having substantial available surface area sufficient to receive deposits of appropriately sized transition metal catalyst.

[0017] The substrate particles are suitably less than about 300 µm. They may be less than 1 µm in transport

bed use. It appears that the transition metal reacts with the substrate crystallites such as to bond the metal to the substrate and fix its position, so as to prevent or retard catalyst agglomeration, at least for so long as it takes to contact the supported transition metal with the suitable carbon source at appropriate reaction conditions. Upon contact, the carbon source seems to pyrolyze on the catalytic site and the desirable morphology fibril grows therefrom.

[0018] As noted, the state of the transition metal catalyst site during fibril formation is important to the practice of this invention. Sometimes, it appears that this desirable catalytic site state as well as the state of the substrate carrier therefore is changing during the whole process hereof. Thus, the catalytic sites may agglomerate or disperse to some extent during the period from introduction into the reaction zone until the fibrils made by the process are recovered. At the time the fibrils are recovered, particles of transition metal catalyst which are sometimes recovered with the fibrils are of 3.5 to 70 nm (35 to 700 Å), preferably 6 to 30 nm (60 to 300 Å) in size. Thus, it is believed that the size of the active catalyst site during fibril formation is substantially comparable to the diameter of the fibril being formed.

[0019] It appears that as fibril formation takes place, active catalyst sites become catalytically expended and need to be replaced. Additionally, it has been found that the fibril forming process is more efficient and capable of better control if the catalyst is added to the reaction zone intermittently or continuously over substantially the entire course of the reaction, or at least a substantial portion thereof. It is possible that the catalyst containing substrate of this invention may ablate with use. That is, when a fibril is formed on a particular catalytic site, that fibril and its associated site may break off from the substrate, with or without some of the substrate, thereby exposing further catalytic sites which were previously inside the substrate particle. Thus, periodic or continuous addition of fresh catalyst is desirable.

[0020] Thus, according to this invention, the fibril forming process hereof is preferably substantially continuous in that a suitable source of carbon, with or without carrier gas, and catalyst containing particles are continuously or intermittently fed to a reaction zone maintained at a fibril forming temperature appropriate to the carbon source being used; while fibril product, usually admixed with the remnants of the catalyst and sometimes substrate as well, are continuously or intermittently recovered.

[0021] The transition metals may be deposited on the substrate by any commonly used technique for accomplishing such deposition. Vapor deposition, sputtering and impregnation may all be suitable. In particular, it has been found to be expeditious to form a water solution or dispersion of the desired metal or metals, mix the water phase with appropriately sized substrate, and then precipitate the metal(s) onto the substrate, e.g. by evaporating the water or any other conventional means.

[0022] It is also possible to deposit the desired transition metals from an organic (as opposed to aqueous) medium. Suitably the transition metals can be dissolved or suspended in such medium, for example, as an organometallic compound, and then impregnated onto and into a suitable substrate. The organic carrier medium is removed, leaving behind the impregnated, deposited transition metals.

[0023] After the transition metals are combined with the substrate as aforesaid, it may be important to treat this combination so as to activate it for this particular catalytic purpose, e.g., by heating it to separate the metals from other ligands, if any, in the deposition compound. It may also be necessary to adjust the size of the prepared catalyst to make it suitable for use in this invention. Commminution or agglomeration, e.g. by binding, may be desirable to produce particles of the proper size, i.e. of less than about 400 μm .

[0024] The catalyst for use in this invention may be put on the substrate hereof in any form or chemical oxidation state. It may be the oxide or have some other ligand. It may be reduced prior to use, but this is not necessary since the fibril forming reaction is a reducing environment and thus the transition metal will be reduced during, or immediately prior to, fibril forming use.

[0025] Fibrils which are very thin and long, diameters of 3.5 to 70 nm and L/D of up to 100 or more, are produced using these catalysts. These fibrils, as produced by this process, without the necessity of further treatment, and without the coproduction of a thermal carbon overcoat, comprise a carbon layer generally concentric about an axis which comprises multiple essentially continuous layers of ordered carbon atoms, which preferably and usually are crystalline and graphitic. This, as produced, outer layer of ordered carbon atoms often surrounds an inner layer of less ordered carbon atoms. Most preferred products of this invention are high yields of high quality, thin fibrils of appropriate long length having substantially uniform, concentric, substantially continuous, ordered, multiple layers of carbon about an axial (inner core) region, which as a different composition/crystallinity and is preferably hollow. Such fibrils preferably have up to about 100 times, and more greater length than diameter, have diameters of up to 70nm (700 angstroms) and are substantially cylindrical about a substantially hollow core and graphitic as made and without having been treated at higher temperatures than the original fibril manufacturing temperature.

[0026] According to one aspect of this invention, operating with catalyst particles as herein set forth, yields of fibrils of greater than about 30 times the weight of transition metal in the catalyst are achievable. In many cases, particularly with mixed transition metals, yields of between 100 and 200 times the weight of transition metal in the catalyst have been achieved. It has been found that in comparable processes, combinations of transition metal catalysts have sometimes increased

yields by a factor of as much as 2 or even more.

[0027] The following examples illustrate the practice of this invention. By following one or more of these examples, high yields of unique fibrils as above described are produced.

Example 1 (comparative)

[0028] A catalyst was prepared using Degussa fumed alumina with an average particle size of about 10nm (100A) and an aggregate mesh size of -100 (149 μm). Iron acetylacetonate was deposited on these alumina particles in a ratio of about 1 part iron, as the acetylacetonate, to 10 parts by weight of alumina. The resultant particle was heated under a hydrogen/ethylene atmosphere under reaction conditions.

[0029] A one (1) inch tube was heated to about 650°C while it was being purged with argon. A mixed flow of hydrogen, at 100 ml/min, and ethylene, at 200 ml/min, was fed to the hot tube for five minutes whereupon catalyst was introduced into the reactor tube. The ethylene/hydrogen mixture was continued through the tubular reactor for 0.5 hours after which the reactor was allowed to cool to room temperature under argon. Harvesting of the fibrils so produced showed a yield of greater than 30 times the weight of the iron in the catalyst.

Example 2

[0030] Into a 3 L. round bottom flask was added 80.08 g of Degussa fumed alumina and 285 ml of methanol. The mixture was stirred to produce a thick paste before a solution of 78.26 g (0.194 moles) of ferric nitrate nonahydrate and 4.00 g (0.0123 moles) of molybdenum(VI) oxide bis(2,4-pentanedionate) in 300 ml of methanol (Fe to Mo atom ratio of 94:6) was added slowly. The thick paste which had collected on the sides of the flask was washed down with 65 ml of additional methanol and the mixture was stirred for 1 hour before house vacuum of 1.1 bar (28 in. Hg) was applied while stirring overnight. The purple-tinted solid was placed in a vacuum of 1.1 bar (28 in. Hg) oven at 100°C for 29 hours. A total of 100.7 g of catalyst was obtained. The catalyst was ground and passed through an 80 mesh (177 μm) sieve prior to use. Analysis of the catalyst indicated 9.43% by weight iron and 0.99% by weight molybdenum.

[0031] A vertical furnace containing a 2.54cm (1 inch) quartz tube with an internal quartz wool plug and thermocouple was equilibrated at 650°C under a down flow of 100 ml/min. hydrogen and 200 ml/min. ethylene. Into the tube (onto the quartz wool plug) was added 0.1044 g of the above-described catalyst. After 30 min., the hydrogen/ethylene flow was stopped and the oven was allowed to cool to near room temperature. A total of 1.2434 g of fibrils was harvested for a yield ratio of 126 times the iron weight content of the catalyst.

Example 3

[0032] A sample of catalyst from example 2 (1.6371 g) was placed in a horizontal furnace under argon and was heated to 300°C. After 30 min. at this temperature, the furnace was cooled and 1.4460 g of catalyst was recovered (12% wt. loss), having 11.1% by weight iron and 1.2% by weight molybdenum.

[0033] A vertical tube furnace containing a 2.54 cm (1 in.) quartz tube with an internal quartz wool plug and thermocouple was equilibrated at 650°C under a 100 ml/min. down flow of hydrogen and 200 ml/min. down flow of ethylene. Into the hot tube was added 0.1029 g of the catalyst described above. After 30 min., the hydrogen/ethylene flow was stopped and the oven was allowed to cool to near room temperature under argon. A total of 1.3750 g of fibrils was isolated for a weight yield based on theoretical iron content of 120 times the iron content.

Example 4

[0034] The vertical tube furnace described in Example 2 was equilibrated at 700°C under the flow of 100 ml/min. hydrogen and 200 ml/min. propane. Onto the quartz wool plug was added 0.1041g of catalyst from Example 2. After 30 min. the fuel gases were stopped and the product was cooled under argon. A total of 0.3993 of fibrils was isolated for a weight yield of 41 times the catalyst iron content.

Example 5

[0035] The procedure of Example 4 was followed at 650°C using 0.1004 g of catalyst from Example 2. A total of 0.3179 g of fibrils was harvested for a weight yield of 34 times the iron content of the catalyst.

Example 6

[0036] Into a round bottom flask was added 4.25 g of Degussa fumed alumina and 30 ml of methanol. The mixture was mechanically stirred while a solution of 4.33 g (10.7 mmol) of ferric nitrate nonahydrate and 0.51 g (1.56 mmol) of molybdenum(VI)oxide bis(2, 4-pentanedionate) in 50 ml of methanol was slowly added. The mixture was stirred for 1 hour before the solvent was removed with the aid of a rotary evaporator. The resulting damp solid was vacuum dried at 105°C, 1.1 bar (28 in. Hg) for 18 hours. The resulting catalyst was ground and passed through an 177µm (80 mesh) sieve. A total of 5.10 g of catalyst was obtained. Analysis of the catalyst indicated 9.04% by weight iron and 2.18% by weight molybdenum to be present.

[0037] Fibrils were prepared following the procedure of Example 2 at 650°C using 0.0936 g of the above catalyst. A total of 0.9487 g of fibrils was isolated for a weight yield of 126 times the catalyst iron content.

Example 7

[0038] Into a round bottom flask was added 3.80 g of Degussa fumed alumina and 30 ml of methanol. The mixture was mechanically stirred while a solution of 4.33 g (10.7 mmol) of ferric nitrate nonahydrate and 2.04 g (6.25 mmol) of molybdenum(VI)oxide bis(2, 4-pentanedionate) in 100 ml of solvent was added. The mixture was held at 105°C and 1.1 bar (28 in. Hg) for 17 hrs. The dried catalyst was sieved (177µm(80 mesh)) to produce 6.10 g of powder. Analysis of the catalyst indicated 8.61% iron and 8.13% molybdenum by weight.

[0039] Fibrils were prepared following the procedure of Example 2 at 650°C using 0.1000 g of the above catalyst. A total of 0.8816 g of fibrils was isolated for a weight yield of 102 times the catalyst iron content.

Example 8

[0040] The procedure of Example 7 was followed at 700°C using methane and 0.1016g of catalyst. A total of 0.0717g of fibrils was isolated for a yield of 8.2 times the iron content of the catalyst.

Example 9

[0041] Into a 500 ml round bottom flask was placed 4.37 g of Degussa fumed alumina and 28 ml of methanol. To the stirred mixture was added a solution of 4.33 g (10.7 mmol) of ferric nitrate nonahydrate and 0.46 g (1.32 mmol) of chromium acetylacetonate in 75 ml of methanol. The mixture was stirred for 1 hr. before it was dried for 18 hr. at 105°C and 1.1 bar (28 in. Hg). The catalyst was ground and sieved (177µm(80 mesh)) to produce 5.57 g of powder. The theoretical metal content by weight was 11.9% iron and 1.4% chromium.

[0042] Fibrils were prepared following the procedure of Example 2 at 650°C using 0.0976 g of the above catalyst. A total of 0.9487 g of fibrils was isolated for a yield of 82 times the theoretical iron content.

Example 10 (comparative)

[0043] Into a 500 ml round bottom flask was placed 4.40 g of Degussa fumed alumina and 35 ml of methanol. To the thick paste was added 4.32g (10.7 mmol) of ferric nitrate nonahydrate in 35 ml of methanol. The mixture was stirred for 45 min. before the solid was dried at 95°C and 1.1 bar (28 in. Hg) for 18 hr. The catalyst was ground and sieved(177µm(80 mesh)).

[0044] Fibrils were prepared following the procedure of Example 2 at 650°C using 0.0930 g of the above catalyst. A total of 0.4890 g of fibrils was isolated for a weight yield of 46 times the catalyst iron content.

Example 11(comparative)

[0045] Into a round bottom flask was placed 4.33 g of

Degussa fumed alumina in 30 ml of methanol. To the stirred paste was added a solution of 4.33 g (10.7 mmol) of ferric nitrate nonahydrate and 0.42 g (1.19 mmol) of ferric acetylacetonate in 50 ml of methanol. The mixture was stirred for 75 min. before drying at 105°C and 1.1 bar (28 in. Hg) for 17 hrs. The solid was ground and sieved (177µm (80 mesh)) to yield 5.87 g of catalyst. Analysis showed 13.79% iron present in the catalyst.

[0046] Fibrils were prepared following the procedure of Example 2 at 650°C using 0.0939 g of the above catalyst to produce 0.3962 g of fibrils. This corresponds to 31 times the iron content of the catalyst.

Example 12

[0047] Into a round bottom flask was added 4.33g of Degussa fumed alumina in 20 ml of water followed by a solution of 4.33 g (10.7 mmol) of ferric nitrate nonahydrate and 0.17g (.138 mmol) of ammonium molybdate in 40 ml of water. The mixture was mechanically stirred for 1 hour. The water was removed at reduced pressure at 40°C overnight. Final drying was accomplished at 140°C and 0.039 bar (26 mm. Hg) for 21 hours to produce 5.57 g of solid. Analysis of the catalyst showed 9.87% by weight iron and 1.45% by weight molybdenum to be present.

[0048] Fibrils were prepared following the procedure of Example 2 at 650°C using 0.0794 g of catalyst to produce 0.8656g of fibrils. This corresponds to 111 times the iron content of the catalyst.

Example 13

[0049] Into a round bottom flask, containing 4.33 g of Degussa fumed alumina and 30 ml of methanol, was added a solution of 4.33 g (10.7 mmol) of ferric nitrate nonahydrate and 0.16 g (0.368 mmol) of ceric nitrate in 50 ml of methanol. An additional 20 ml of methanol was used to wash all the salts into the flask. The mixture was stirred for one hour before the solvent was removed at reduced pressure. The solid was dried at 130°C and 0.040 bar (27 mm Hg) for four days to produce 5.32 grams of catalyst. Analysis of the solid indicated 9.40% iron and 0.89% cerium to be present.

[0050] Fibrils were prepared following the procedure of Example 2 at 650°C using 0.0941g of catalyst to produce 0.7552g of fibrils. This corresponds to 88 times the iron content of the catalyst.

Example 14

[0051] Into a round bottom flask was added 4.33g of Degussa fumed alumina and 30 ml of methanol. Onto the alumina was poured a solution of 4.33g (10.7 mmol) of ferric nitrate and 0.31g (1.22 mmol) of manganese(II) acetylacetonate in 50 ml of methanol. The solvent was removed at reduced pressure of 0.040 bar (27 mm Hg) and the damp solid was vacuum dried at 140°C to pro-

duce 5.18g of solid. Analysis of the catalyst indicated 9.97% iron and 1.18% manganese.

[0052] Fibrils were prepared following the procedure of Example 2 at 650°C using 0.070g of catalyst to produce 0.4948g of fibrils. This corresponds to 66 times the iron content of the catalyst.

Example 15

[0053] Into a round bottom flask was added 4.33g of Degussa fumed alumina and 30 ml of methanol. Onto the alumina was poured a solution of 4.33g (10.7 mmol) of ferric nitrate and 0.43g (1.22 mmol) of manganese(II) acetylacetonate in 50 ml of methanol. The solvent was removed at reduced pressure and the damp solid was vacuum dried at 140°C to produce 5.27g of solid. Analysis of the catalyst indicated 10.00% iron and 1.18% manganese by weight.

[0054] Fibrils were prepared following the procedure of Example 2 at 650°C using 0.0723g of catalyst to produce 0.7891g of fibrils. This corresponds to 110 times the iron content of the catalyst on a weight basis.

Example 16 (comparative)

[0055] Degussa fumed alumina (400g) and deionized water (8.0L) were added to a 22 L flask equipped with a stirrer, pH meter and probe, and two 2 L addition funnels. One funnel contained an aqueous solution of ferric nitrate nonahydrate (511g dissolved in 5654 ml of water) and the other an aqueous solution of sodium bicarbonate (480g dissolved in 5700 ml of water).

[0056] The pH of the alumina slurry was first adjusted to 6.0 by adding the sodium bicarbonate solution to raise it or the ferric nitrate solution to lower it. Next, both solutions were added simultaneously over 3-4 hours with good agitation while maintaining the pH at 6.0. When the addition was complete, stirring was continued for an additional 1/2 hour, after which the slurry was filtered on a 32 cm Buchner funnel. The filter cake was then washed with deionized water and returned to the 22 L flask. Next, additional deionized water was added and the slurry stirred for another 1/2 hour. The batch was then filtered, washed with deionized water, and vacuum-dried at 100°C to constant weight (475g). Following drying, the final catalyst was prepared by grinding and sieving the product to 177µm (-80 mesh).

Example 17

[0057] This Example illustrates the practice of this invention using periodic addition of catalyst to produce high fibril yields. A 10.16 cm (four-inch) quartz tube, closed on the bottom, was placed in a 10.16(4 inch)diameter x 60.96 cm (24 inch) long furnace. The tube was purged with argon while being heated to 620°C. When the tube was hot, the gas feed was switched to a mixture of hydrogen (1.0 l/min) and ethyl-

ene (5.6 l/min) via a dip tube to the bottom of the 10.16 cm (4 inch) tube. After 5 min of purging, the catalyst addition was begun.

[0058] A total of 41.13g of catalyst, prepared as described in the Example 16, was added to the hot reactor reservoir. The catalyst was added periodically to the hot reactor in small portions (0.2g) over a period of approximately 6 hours. After catalyst addition was complete, the reaction was allowed to run for an additional one hour and the reactor then cooled to room temperature under argon. The fibrils were removed from the tube and weighed. This batch gave 430g total yield of fibrils which is unusually high for a catalyst based upon iron has the only transition metal. In single batch addition of an iron only catalyst, fibril yields of about 30 times the iron content have been observed whereas here the fibril yield is more than 70 times the iron content of the catalyst.

Example 18

[0059] The tube and furnace described in Example 17 were heated to 650° under an argon purge. When the tube was hot the gas feed was switched to hydrogen and ethylene as described in Example 17.

[0060] A total of 20.4g of catalyst (Fe-Mo) prepared as described in Example 2 was added in a manner similar to that described in Example 17. This batch gave a total fibril yield of 255g.

Claims

1. A continuous process for producing carbon fibrils by decomposing a source of carbon at elevated temperatures in contact with a multivalent metal and recovering the fibrils formed thereby comprising the steps of introducing catalyst particles having a size of up to 400 μm and comprising, as the multivalent metal iron mixed with at least one other transition metal on a particulate substrate into a reactor heated to a temperature of 500°C to 1500°C, and recovering the fibrils formed thereby wherein said metal is present on said substrate as a multiplicity of discontinuous catalytic sites which, at least during fibril formation, have a size of 3.5 to 70 nm (35 to 700 Å) which size is measured by measuring the size of the transition metal particles recovered along with produced fibrils.
2. The process of claim 1 wherein said transition metal comprises iron with a catalyst site size of about 6 to 30 nm (60 to 300 Å).
3. The process of claim 1 wherein said other transition metal is at least one selected from the group consisting of molybdenum and chromium.
4. The process of claim 1 wherein said carbon source is a mixture of hydrocarbons.
5. The process of claim 1 wherein said carbon source is at least one member selected from the group consisting of methane, ethane, propane, butane, benzene, butene, isobutene, cyclohexane, ethylene, propylene, acetylene, toluene, xylene, cumene, ethyl benzene, naphthalene, phenanthrene, anthracene, formaldehyde, acetaldehyde, acetone, methanol, ethanol, and carbon monoxide.
6. The process as claimed in claim 1 carried out in an ebulliating bed wherein said catalyst particles have a particle size of about 50 to 300 μm (microns).
7. The process of claim 1 carried out in a transport bed wherein said catalyst particles have a particle size of less than about 10 μm (microns).
8. The process of claim 7 wherein said catalyst particles have a particle size of less than about 1 μm (micron).
9. The process of claim 1 including physically cofeeding said catalyst particles and said source of carbon downflow through a reaction zone.
10. The process of claim 1 including physically cofeeding said catalyst particles and said source of carbon upflow through a reaction zone.
11. The process of claim 1 including cofeeding a non-hydrocarbonaceous gas with said carbon contributing feed.
12. The process of claim 1 wherein said catalyst particles are added to the heated reactor over the course of the reaction.
13. The process of claim 12 wherein said catalyst particles are added intermittently over the course of the reaction.
14. The process of claim 1 wherein said source or carbon is added to the heated reactor.
15. The process of claim 1 wherein said source of carbon is added to the heated reactor over the course of the reaction.
16. The process of claim 1 wherein said catalyst particles and said source of carbon are added to the heated reactor over the course of the reaction.
17. The process of claim 1 including recovering a high yield of product comprising carbon fibrils characterized as tubes that are substantially free of pyrolytically deposited thermal carbon and have graphitic

layers that are substantially parallel to the fibril axis.

18. The process of claim 1 wherein the diameter of said fibrils is between 3.5 and 70 nm, inclusive.

19. The process of claim 1 wherein said process is carried out in a fluidized bed.

20. A process for continuously producing high yields of long, thin fibrils comprising at least long portions having atomically ordered outer surfaces, as made by contacting a vaporous source of the atoms which will comprise said outer surface with catalyst particles; which process comprises introducing said catalyst particles into a reactor heated to a temperature of 500°C to 1500°C, said catalyst particles having a size of up to 400 µm (microns) and comprising, as the multivalent metal iron mixed with at least one other transition metal on a particulate substrate, said metal being present on said substrate as a multiplicity of discontinuous catalytic sites which, at least during fibril formation, have a size of 3.5 to 70 nm (35 to 700 Å) which size is measured by measuring the size of the transition metal particles recovered along with produced fibrils, and recovering the fibrils formed thereby.

Patentansprüche

1. Kontinuierliches Verfahren zur Erzeugung von Kohlenstoff-Fibrillen durch Zersetzen einer Kohlenstoff-Quelle bei erhöhter Temperatur in Gegenwart eines multivalenten Metalls und Wiedergewinnen der dadurch gebildeten Fibrillen, umfassend das Einführen von Katalysator-Teilchen mit einer Größe bis zu 400 µm und umfassend Eisen als multivalentes Metall, das mit zumindest einem anderem Übergangsmetall gemischt ist, auf einem körnigen Substrat in einen Reaktor, der auf eine Temperatur von 500 bis 1500°C erwärmt ist, und Wiedergewinnen der dadurch gebildeten Fibrillen, worin das Metall auf dem Substrat als eine Vielzahl von diskontinuierlichen, katalytischen Stellen vorhanden ist, die zumindest während der Fibrillenbildung eine Größe von 3,5 bis 70 nm (35 bis 700 Å) haben, wobei die Größe durch Messen der Größe der Übergangsmetall-Teilchen gemessen wird, die zusammen mit erzeugten Fibrillen wiedergewonnen werden.
2. Verfahren nach Anspruch 1, worin das Übergangsmetall Eisen mit einer Größe der Katalysatorstelle von etwa 6 bis 30 nm (60 bis 300 Å) umfaßt.
3. Verfahren nach Anspruch 1, worin das andere Übergangsmetall zumindest eines ist, ausgewählt aus der Gruppe, bestehend aus Molybdän und Chrom.

4. Verfahren nach Anspruch 1, worin die Kohlenstoff-Quelle eine Mischung aus Kohlenwasserstoffen ist.
5. Verfahren nach Anspruch 1, worin die Kohlenstoff-Quelle zumindest eine Verbindung ist, ausgewählt aus der Gruppe, bestehend aus Methan, Ethan, Propan, Butan, Benzol, Buten, Isobuten, Cyclohexan, Ethylen, Propylen, Acetylen, Toluol, Xylol, Cumol, Ethylbenzol, Naphthalin, Phenanthren, Anthracen, Formaldehyd, Acetaldehyd, Aceton, Methanol, Ethanol und Kohlenmonoxid.
6. Verfahren nach Anspruch 1, das in einem Siedebett durchgeführt wird, worin die Katalysator-Teilchen eine Teilchengröße von etwa 50 bis 300 µm (Mikron) haben.
7. Verfahren nach Anspruch 1, durchgeführt in einem Transportbett, worin die Katalysator-Teilchen eine Teilchengröße von weniger als etwa 10 µm (Mikron) haben.
8. Verfahren nach Anspruch 7, worin die Katalysator-Teilchen eine Teilchengröße von weniger als etwa 1 µm (Mikron) haben.
9. Verfahren nach Anspruch 1, umfassend das physikalische gemeinsame Führen der Katalysator-Teilchen und der Kohlenstoff-Quelle im Abwärtsstrom durch eine Reaktionszone.
10. Verfahren nach Anspruch 1, umfassend das gemeinsame physikalische Führen der Katalysator-Teilchen und der Kohlenstoff-Quelle im Aufwärtsstrom durch eine Reaktionszone.
11. Verfahren nach Anspruch 1, umfassend das gemeinsame Führen eines nicht-Kohlenwasserstoff-haltigen Gases mit der Kohlenstoff-haltigen Zufuhr.
12. Verfahren nach Anspruch 1, worin die Katalysator-Teilchen zu dem erwärmten Reaktor während der Reaktion zugegeben werden.
13. Verfahren nach Anspruch 12, worin die Katalysator-Teilchen absatzweise während des Verlaufes der Reaktion zugegeben werden.
14. Verfahren nach Anspruch 1, worin die Kohlenstoff-Quelle zu dem Heizreaktor gegeben wird.
15. Verfahren nach Anspruch 1, worin die Kohlenstoff-Quelle zu dem erwärmten Reaktor während des Verlaufes der Reaktion gegeben wird.
16. Verfahren nach Anspruch 1, worin die Katalysator-Teilchen und die Kohlenstoff-Quelle zu dem Heizre-

aktor während des Verlaufes der Reaktion gegeben werden.

17. Verfahren nach Anspruch 1, umfassend das Wiedergewinnen einer hohen Ausbeute des Produktes, umfassend Kohlenstoff-Fibrillen, gekennzeichnet als Rohre, die im wesentlichen frei sind von pyrolytisch niedergeschlagenem, thermischem Kohlenstoff und graphitische Schichten haben, die im wesentlichen parallel zu der Fibrillachse sind.

18. Verfahren nach Anspruch 1, worin der Durchmesser der Fibrillen zwischen 3,5 und 70 nm einschließlich ist.

19. Verfahren nach Anspruch 1, worin das Verfahren in einem Fließbett durchgeführt wird.

20. Verfahren zum kontinuierlichen Erzeugen hoher Ausbeuten von langen, dünnen Fibrillen, umfassend zumindest lange Bereiche mit atomisch angeordneten äußeren Oberflächen, erzeugt durch Kontaktieren einer dampfartigen Quelle der Atome, die die äußere Oberfläche umfassen, mit Katalysator-Teilchen; wobei das Verfahren das Einführen von Katalysator-Teilchen in einen Reaktor, der auf eine Temperatur von 500 bis 1500°C erwärmt ist, wobei die Katalysator-Teilchen eine Größe von bis zu 400 µm (Mikron) haben und als multivalentes Metall Eisen, das mit zumindest einem anderen Übergangsmetall vermischt ist, auf einen körnigen Substrat enthalten, wobei das Metall auf dem Substrat als eine Vielzahl von diskontinuierlichen katalytischen Stellen vorhanden ist, die zumindest während der Fibrill-Bildung eine Größe von 3,5 bis 70 nm (35 bis 700 Å) haben, wobei die Größe durch Messen der Größe der Übergangsmetall-Teilchen gemessen wird, die zusammen mit den erzeugten Fibrillen wiedergewonnen werden, und das Wiedergewinnen der dadurch gebildeten Fibrillen umfaßt.

Revendications

1. Procédé en continu destiné à la production de fibrilles de carbone par décomposition d'une source de carbone à des températures élevées en contact avec un métal plurivalent et récupération des fibrilles ainsi formées, comprenant les étapes consistant à introduire des particules de catalyseur présentant une dimension allant jusqu'à 400 µm et comprenant, en tant que métal plurivalent, du fer mélangé à au moins un autre métal de transition sur un substrat particulaire dans un réacteur chauffé à une température de 500° C à 1 500° C, et à récupérer les fibrilles ainsi formées, dans lequel ledit métal est présent sur ledit substrat sous forme d'une multiplicité de sites catalytiques discontinus lesquels, au moins pendant la formation des

fibrilles, présentent une dimension de 3,5 à 70 nm (35 à 700 Å), laquelle dimension est mesurée en mesurant la dimension des particules de métal de transition récupérées en même temps que les fibrilles produites.

2. Procédé selon la revendication 1, dans lequel ledit métal de transition comprend du fer présentant une dimension de sites de catalyseur d'environ 6 à 30 nm (60 à 300 Å).

3. Procédé selon la revendication 1, dans lequel ledit autre métal de transition est au moins l'un choisi parmi le groupe constitué du molybdène et du chrome.

4. Procédé selon la revendication 1, dans lequel ladite source de carbone est un mélange d'hydrocarbures.

5. Procédé selon la revendication 1, dans lequel ladite source de carbone est au moins un élément choisi parmi le groupe constitué du méthane, de l'éthane, du propane, du butane, du benzène, du butène, de l'isobutène, du cyclohexane, de l'éthylène, du propylène, de l'acétylène, du toluène, du xylène, du cumène, de l'éthylbenzène, du naphthalène, du phénanthrène, de l'anthracène, du formaldéhyde, de l'acétaldéhyde, de l'acétone, du méthanol, de l'éthanol, et du monoxyde de carbone.

6. Procédé selon la revendication 1, exécuté dans un lit à ébullition dans lequel lesdites particules de catalyseur présentent une dimension de particules d'environ 50 à 300 µm (microns).

7. Procédé selon la revendication 1, exécuté dans un lit à transport dans lequel lesdites particules de catalyseur présentent une dimension de particules de moins d'environ 10 µm (microns).

8. Procédé selon la revendication 7, dans lequel lesdites particules de catalyseur présentent une dimension de particules de moins d'environ 1 µm (microns).

9. Procédé selon la revendication 1, comprenant une coalimentation physique desdites particules de catalyseur et de ladite source de carbone en circulation vers le bas à travers une zone réactionnelle.

10. Procédé selon la revendication 1, comprenant une coalimentation physique desdites particules de catalyseur et de ladite source de carbone en circulation vers le haut à travers une zone réactionnelle.

11. Procédé selon la revendication 1, comprenant une coalimentation d'un gaz non-hydrocarboné avec

ladite alimentation de contribution en carbone.

12. Procédé selon la revendication 1, dans lequel lesdites particules de catalyseur sont ajoutées au réacteur chauffé pendant la durée de la réaction. 5
13. Procédé selon la revendication 12, dans lequel lesdites particules de catalyseur sont ajoutées par intermittence pendant la durée de la réaction. 10
14. Procédé selon la revendication 1, dans lequel ladite source de carbone est ajoutée au réacteur chauffé. 15
15. Procédé selon la revendication 1, dans lequel ladite source de carbone est ajoutée au réacteur chauffé pendant la durée de la réaction. 20
16. Procédé selon la revendication 1, dans lequel lesdites particules de catalyseur et ladite source de carbone sont ajoutées au réacteur chauffé pendant la durée de la réaction. 25
17. Procédé selon la revendication 1, comprenant la récupération d'un rendement élevé de produit comprenant des fibrilles de carbone caractérisées comme étant des tubes qui sont pratiquement exempts de carbone thermique déposé de façon pyrolytique et comportent des couches graphitiques qui sont pratiquement parallèles à l'axe des fibrilles. 30
18. Procédé selon la revendication 1, dans lequel le diamètre desdites fibrilles est entre 3,5 et 70 nm, inclus. 35
19. Procédé selon la revendication 1, dans lequel ledit procédé est exécuté dans un lit fluidisé. 40
20. Procédé destiné à produire en continu des rendements élevés de fibrilles longues, fines, comprenant au moins des parties longues présentant des surfaces externes atomiquement ordonnées, réalisées en mettant en contact une source sous forme de vapeur des atomes qui constitueront ladite surface externe avec des particules de catalyseur, lequel procédé comprend l'introduction desdites particules de catalyseur dans un réacteur chauffé à une température de 500° C à 1 500° C, lesdites particules de catalyseur présentant une dimension allant jusqu'à 400 µm (microns) et comprenant, en tant que métal plurivalent, du fer mélangé à au moins un autre métal de transition sur un substrat particulaire, ledit métal étant présent sur ledit substrat sous forme d'une multiplicité de sites catalytiques discontinus lesquels, au moins pendant la formation des fibrilles, présentent une dimension de 3,5 à 70 nm (35 à 700 Å), laquelle dimension est mesurée en mesurant la dimension des particules 55

de métal de transition récupérées en même temps que les fibrilles produites, et la récupération des fibrilles ainsi formées.